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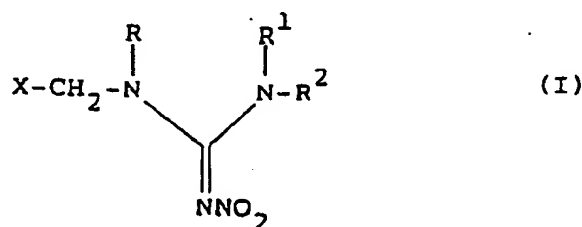
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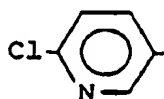
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(54) **Nitroguanidine compounds and insecticides.**

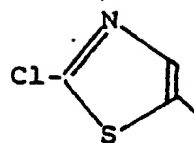
(57) Insecticides which are improved over the conventional insecticides are provided by the use of a nitroguanidine compound represented by the following general formula (I):



wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by



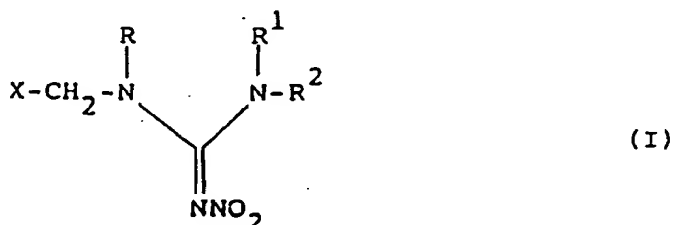
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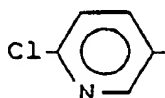
The insecticides provided by this invention are extremely active for controlling various harmful insects and yet are low in toxicity to warm-blooded animals, fishes and Crustacea, the remaining quantity thereof after their use is small, and they do not produce phytotoxicity in various plants.

## NITROGUANIDINE COMPOUNDS AND INSECTICIDES

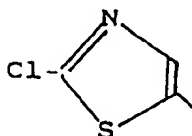
The present invention relates to nitroguanidine compounds, a process for preparing the same and insecticides containing the same as active ingredients. The nitroguanidine compounds provided by this invention are represented by the following general formula (I):



wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by



or



A variety of chemicals have been developed and used as insecticides for controlling harmful insects. Representative examples of such chemicals are organic phosphorous compounds, carbamate compounds and synthetic pyrethroids. However, by the repeated use of these insecticides, many harmful insects have acquired resistance to these chemicals, thus making it difficult to control such insects. On the other hand, although some conventional insecticides have high insecticidal activity, they might cause environmental pollution problems either due to their high toxicity to warm-blooded animals or fishes and Crustacea or due to their remaining in the environment in large amounts for a long time after use, thus putting the ecological system out of order. Accordingly, there is a demand for the development of novel insecticides which more effectively control harmful insects, which have acquired resistance to conventional chemicals, and yet are low in toxicity not only to warm-blooded animals but also fishes and Crustacea. There is also a demand for novel insecticides which remain in plants and soils in decreased amounts after their use, and do not produce phytotoxicity in various plants.

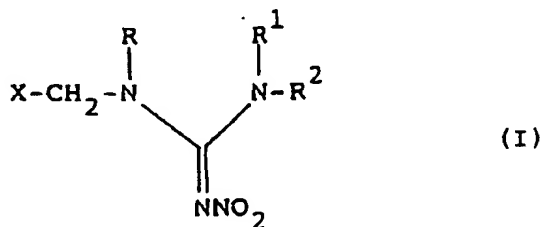
An object of this invention is to provide novel compounds which (1) more effectively control harmful insects that have acquired resistance to conventional chemicals, (2) are low in toxicity not only to warm-blooded animals but also fishes and Crustacea and (3) remain in decreased amounts in plants and soils after their use. Lastly, such novel compounds do not produce phytotoxicity in various plants.

Another object of this invention is to provide a process for preparing such compounds.

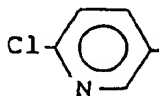
A further object of this invention is to provide improved insecticides which (1) more effectively control harmful insects having resistance to conventional insecticides, (2) are low in toxicity not only to warm-

blooded animals but also fishes and Crustacea, and (3) remain in decreased amounts in plants and soils after their use. Lastly, such improved insecticides do not produce phytotoxicity in various plants.

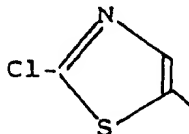
The subject matter of the invention comprises novel nitroguanidine compounds represented by the following general formula (I):



wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by



or

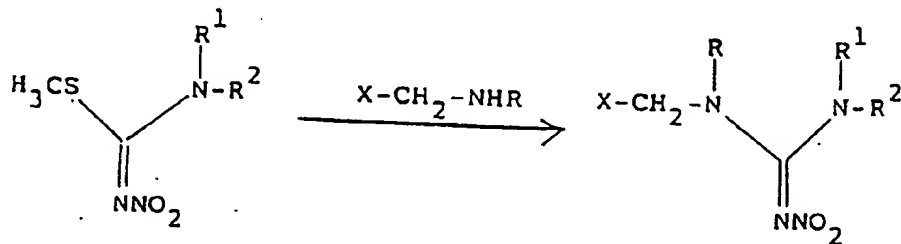


These compounds can be used as extremely powerful insecticides which exhibit rapid effects.

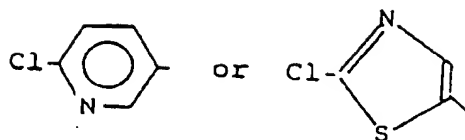
The compounds of this invention represented by the general formula (I) are novel compounds which have not been described in any prior publication.

These compounds may be prepared by the processes described below.

#### Preparation Process (a)



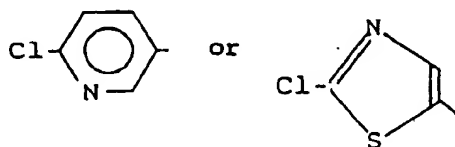
wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by



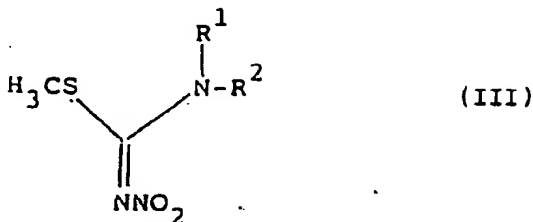
10 In the preparation process (a), as indicated by the reaction formula set forth above, the compounds of this invention may be readily prepared by reacting, with or without solvents a compound represented by the general formula (II):



wherein R stands for hydrogen or methyl, and X stands for a group indicated by



25 with a compound represented by the general formula (III):



wherein  $R^1$  and  $R^2$  are the same or different, and stand for hydrogen or methyl.

40 Examples of the suitable solvents which may be used in this reaction are: ethers such as diethyl ether and tetrahydrofuran; alcohols such as methanol and ethanol; acetonitrile; aromatic hydrocarbons; chlorinated hydrocarbons; DMF; and DMSO. These solvents may be used alone or in combination. Particularly preferable solvents are alcohols such as methanol and ethanol.

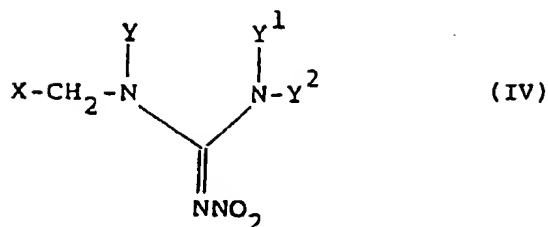
The reaction temperature may be freely set within a range of from room temperature to  $150^\circ\text{C}$ , preferably from room temperature to  $80^\circ\text{C}$ .

#### 45 Preparation Process (b)

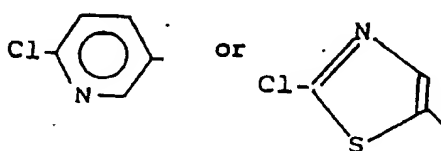
This process may be adopted when the groups R,  $R^1$  and  $R^2$  of general formula (I) are in a certain special interrelation.

50 More particularly, according to this process, the compounds of this invention may be readily prepared by reacting a compound represented by the general formula (IV):

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wherein Y, Y<sup>1</sup> and Y<sup>2</sup> are the same or different, and stand for hydrogen or methyl provided that at least one of them is hydrogen; and X stands for a group indicated by

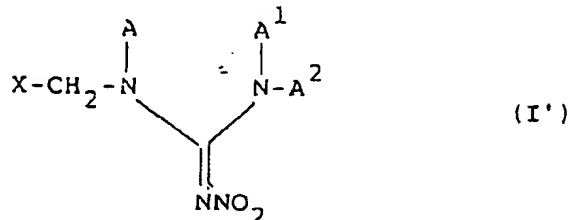


with a compound (V) represented by the general formula (V):

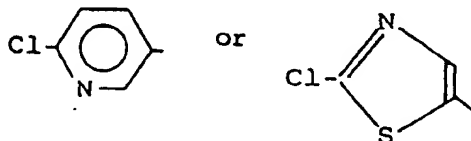


wherein Z stands for halogen;

in the presence of a basic material, generally by using the compound (V) in a molar ratio of from 1 to 3.3 mols per 1 mol of the compound (IV), whereby a compound of this invention represented by the following formula (I') is prepared.



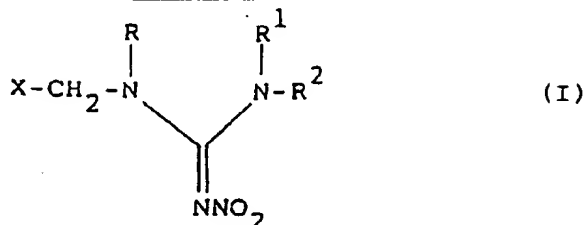
wherein A, A<sup>1</sup> and A<sup>2</sup> are the same or different, and stand for hydrogen or methyl provided that at least one of them is methyl, and X stands for a group indicated by



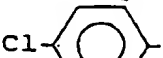
Examples of solvent which may be used in this reaction are: ethers such as diethyl ether and tetrahydrofuran; acetonitrile; aromatic hydrocarbons; chlorinated hydrocarbons; DMF; and DMSO. These solvents may be used alone or in combination. Particularly preferable solvents are DMF and DMSO. Compounds which may be used as the basic materials are, for example, inorganic bases such as sodium hydride, sodium hydroxide and potassium carbonate, and organic bases such as DBU. These bases may be used in a molar ratio of from 1 to 3.3 per 1 mol of the compound (IV). The reaction temperature may be set within a range of from -(minus) 20° C to 100° C, preferably from -(minus) 10° C to 60° C.

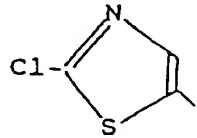
It should be noted again that the compounds of this invention are novel compounds. Typical compounds (I) of the present invention prepared by the above processes are set forth in the following Table 1.

Table 1



Compound No.	R	R <sup>1</sup>	R <sup>2</sup>	Melting Point
1	CH <sub>3</sub>	H	H	Decomposed at 158.0°C
2	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	99.0 to 100.0 °C
3	H	CH <sub>3</sub>	H	152.5 to 153.5 °C
4	H	H	H	198.0 to 199.0 °C
5	CH <sub>3</sub>	CH <sub>3</sub>	H	130.0 to 132.0 °C
6	CH <sub>3</sub>	H	H	116.0 to 117.0 °C
7	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	101.0 to 102.0 °C
8	H	CH <sub>3</sub>	H	151.0 to 153.0 °C
9	H	H	H	160.0 to 162.0 °C

Note: The compounds denoted by 1 to 5 are compounds represented by the general formula (I) wherein X is ; and

the compounds denoted by 6 to 9 are compounds represented by the general formula (I) wherein X is 

An effective amount of any of the compounds of this invention may be applied directly as an insecticide or may be formulated by conventional technology in the form of an emulsion, a wettable powder, a dust, a granule or in a flowable form, and then applied as a formulated product. A liquid or solid carrier may be used for preparing a formulated product containing one or more of the compounds of this invention. Liquid carriers which may be used for this purpose include, for example, organic solvents, and conveniently used

are xylene, chlorobenzene, methylnaphthalene, cyclohexanone, isophorone, alcohols, dimethylformamide and N-methylpyrrolidone. Examples of solid carriers include kaoline, talc, bentonite, diatomaceous earth and clay, and synthetic compounds such as alumina, zeolite and silicates may also be used. In preparation of the formulated products, various adjuvants, such as emulsifiers, dispersants, spreaders, wetting agents and penetrating agents, may be used for providing the products with the desired properties of emulsification, dispersion, suspension and penetration.

The compounds of this invention, which are represented by the general formula (I) set forth above, have strong insecticidal activity against various harmful insects, including Hemiptera, Lepidoptera, Coleoptera, Diptera, Orthoptera and Isoptera, and yet are low in toxicity to human beings and animals and do not produce phytotoxicity in various plants, and thus can be practically used as superior insecticides.

Examples of harmful insects which may be effectively controlled by the compounds of this invention are as follows:

Hemiptera such as *Nilaparvata lugens*, *Laodelphax striatellus*, *Nephotettix cincticeps*, *Pseudococcus comstocki*, *Unaspis yanonensis*, *Myzus persicae*, *Aphis gossypii*, *Lipaphis erysimi*, *Stephanitis nashi*, *Scotinophara lurida* and *Trialeurodes vaporariorum*; Lepidoptera such as *Pieris rapae*, *Spodoptera litura*, *Mamestra brassicae*, *Chilo suppressalis*, *Plutera xylostella*, *Adoxophyes orana*, *Agrotis fucosa*, *Cnaphalocrocis medinalis* and *Ostrinia furnacalis*; Coleoptera such as *Henosepilachna vigintioctopunctata*, *Aulacophora femoralis*, *Phyllotreta striolata*, *Lissorhoptrus oryzophilus*, *Sitophilus zeamais* and *Anomala rufocuprea*; Diptera such as *Musca domestica*, *Hylemia platura*, *Culex pipiens*; Orthoptera such as *Gryllotalpa africana*, *Blattella germanica* and *Locusta migratoria*; and Isoptera such as *Coptotermes formosanus* and *Reticulitermes speratus*.

The present invention will now further be explained by way of the following non-limiting examples.

## Examples

The processes for the preparation of the compounds of this invention are described in detail by referring to some Synthesis Examples.

### Synthesis Example 1

#### Synthesis of N-[(6-chloro-3-pyridinyl)methyl]-N-methyl-N'-nitroguanidine

A mixture of 13.8 g of 6-chloro-N-methyl-3-pyridinemethanamine, 10.8 g of S-methyl-N-nitroisothiurea and 60 ml of ethanol was heated at 50 °C with stirring for 2 hours and then refluxed for additional 5 hours. After cooling, the separated crystal was collected by filtration, washed with a small amount of ethanol, and dried to obtain 14.7 g of the title compound. This compound is No.1 in Table 1. m.p.: 158.0 °C (Decomposed).

### Synthesis Example 2

#### Synthesis of N-[(6-chloro-3-pyridinyl)methyl]-N,N',N'-trimethyl-N'-nitroguanidine

To a mixture of 0.21 g of sodium hydride (60% oily) and 4 ml of N,N-dimethylformamide, 1.22 g of N-[(6-chloro-3-pyridinyl)methyl]-N-methyl-N'-nitroguanidine was added in portions with stirring under cooling with ice. Then, the mixture was stirred at room temperature. This was followed by one-hour additional stirring, and thereafter 0.74 g of methyl iodide was added dropwise. After stirring over night, the reaction mixture was poured into ice-water and the insoluble substance was filtered off. The filtrate was extracted with dichloromethane. The dichloromethane layer was washed with water, dried, concentrated. The residue was purified by silica gel column chromatography (Solvent: Chloroform/Methanol = 50/1) to obtain 0.31 g of the title compound. This compound is No. 2 in Table 1. m.p.: 99.0 - 100.0 °C



Synthesis Example 3

## 5 Synthesis of N-[(2-chloro-5-thiazolyl)methyl]-N-methyl-N'-nitroguanidine

A mixture of 1.50 g of 2-chloro-N-methyl-5-thiazolemethanamine, 1.13 g of S-methyl-N-nitrosothiourea and 6 ml of ethanol was heated under reflux for 6 hours. After cooling, the reaction mixture was concentrated under reduced pressure and the residue was purified by silica gel column chromatography  
 10 (Solvent: Chloroform/Methanol=20/1) to obtain 1.00 g of the title compound as crystals. This compound is No.6 in Table 1. m.p.: 116.0 - 117.0 °C

Synthesis Example 4

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## Synthesis of N-[(2-chloro-5-thiazolyl)methyl]-N,N',N'-trimethyl-N''-nitroguanidine

20 0.15 g of sodium hydride (60% oily) was added in portion to a mixture of 0.83 g of N-[(2-chloro-5-thiazolyl)methyl]-N-methyl-N'-nitroguanidine and 10 ml of N,N-dimethylformamide with stirring under cooling with ice-water. After the addition, the reaction mixture was stirred for additional 1.5 hours. 0.54 g. of methyl iodide was added dropwise to the reaction mixture. After the temperature of the reaction mixture was returned to room temperature, it was stirred over night. The reaction mixture was poured into ice water,  
 25 and then extracted with chloroform. The chloroform layer was washed with water, dried, concentrated, and then purified by silica gel column chromatography (Solvent: Chloroform/Methanol = 50/1 to 10/1) to obtain 0.15 g of the title compound as crystals. This compound is No. 7 in Table 1. m.p.: 101.0 - 102.0 °C

30

Synthesis Example 5

## Synthesis of N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitroguanidine

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A mixture of 0.48 g of 2-chloro-5-thiazolemethanamine, 0.31 g of 5-methyl-N-methyl-N'-nitrosothiourea prepared by the method described in the Journal of Medical Chemistry., 1977, Vol. 20, No. 7, P. 905 and 3 ml of ethanol was heated under reflux for 6 hours. After cooling, the reaction mixture was concentrated under reduced pressure, and the residue was purified by silica gel column chromatography (Solvent:  
 40 Chloroform/Methanol = 20/1) to obtain 0.37 g of the title compound as crystals. This compound is No. 8 in Table 1. m.p.: 151.0 - 153.0 °C

45

Synthesis Example 6

## Synthesis of N-[(2-chloro-5-thiazolyl)methyl]-N'-nitroguanidine

50 A mixture of 0.35 g of 2-chloro-5-thiazolemethanamine, 0.31 g of S-methyl-N-nitrosothiourea and 3 ml of ethanol was heated under reflux for 6 hours. After cooling, the separated crystals were collected by filtration, washed with a small amount of ethanol, and then recrystallized from ethanol to obtain 0.42 g of the title compound as crystals. This compound is No. 9 in Table 1. m.p.: 160.0 - 162.0 °C

Some specific Formulation Examples are shown below. However, it is noted that the carriers, surfactants and other additives which may be used in formulation of the insecticides according to this invention are not limited by the following Formulation Examples. In the following Formulation Examples, "part" stands for "part by weight".  
 55

## EP 0 425 978 A2

### Formulation Example 1

32.5 Parts of the Compound No. 1 in Table 1, 3 parts of lignin sulfonic acid, 4 parts of polyoxy thyl ne  
5 alkylphenyl ether, 2 parts of silicon dioxide hydrate and 58.5 parts of clay are well mixed while being  
powdered to obtain a wettable powder.

### Formulation Example 2

10

5.4 Parts of the Compound No. 2 in Table 1, 2 parts of silicon dioxide hydrate and 92.6 parts of talc are  
well mixed while being powdered to obtain a dust.

15

### Formulation Example 3

5.4 Parts of the Compound No. 8 in Table 1, 3 parts of lignin sulfonate, 1 part of sodium dodecylsul-  
20 fonate, 30 parts of bentonite and 60.6 parts of clay are well mixed while being powdered. The mixture is  
then kneaded with water, granulated and dried to obtain granules.

The insecticidal effects of the compounds of this invention are shown by referring to some Test  
Examples.

25

### Test Example 1

Five rice seedlings having a height of 7 cm were dipped, for 10 seconds, into aqueous dilutions of  
30 wettable powder each containing a predetermined concentration of each sample compound, prepared in  
accordance with the procedure as described in Formulation Example 1. After air drying, the roots of the five  
rice seedlings were wrapped with sanitary cotton dampened with water, and placed in a glass cylinder  
having a diameter of 3 cm and a height of 20 cm. Ten second instar larvae of a green rice leafhopper which  
had acquired resistance to chemicals were released into each glass cylinder. The cylinder was allowed to  
35 stand in a room maintained at 26 °C. The mortality of the larvae was examined 48 hours after treatment.  
The results are shown in Table 2.

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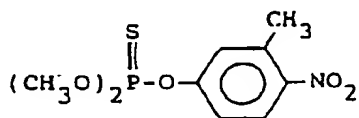
Table 2

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	Compound No.	Mortality (%), 500 ppm
10	1	100
	2	100
	3	100
15	4	100
	5	100
	6	100
20	7	100
	8	100
	9	100
25	Comparative Compound	70

30 Note: The comparative compound set forth in Table 2 is Sumithion  
(Trade Name), represented by the following formula.

35



40

## Test Example 2

45

50 Five rice seedlings having a height of 7 cm were dipped, for 10 seconds, into aqueous dilutions of wettable powder each containing a predetermined concentration of each sample compound, prepared in accordance with the procedure as described in Formulation Example 1. After air drying, the roots of the five rice seedlings were wrapped with sanitary cotton dampened with water, and placed in a glass cylinder having a diameter of 3 cm and a height of 20 cm. Ten second instar larvae of a brown rice planthopper which had acquired resistance to chemicals were released into each glass cylinder. The cylinder was allowed to stand in a room maintained at 26 °C. The mortality of the larvae was examined 48 hours after treatment. The results are shown in Table 3.

55

Table 3

Compound No.	Mortality (%), 500 ppm
1	100
2	100
3	100
4	100
5	100
6	100
7	100
8	100
9	100
Comparative Compound	65

Note: The comparative compound set forth in Table 3 is the same as that in Table 2.

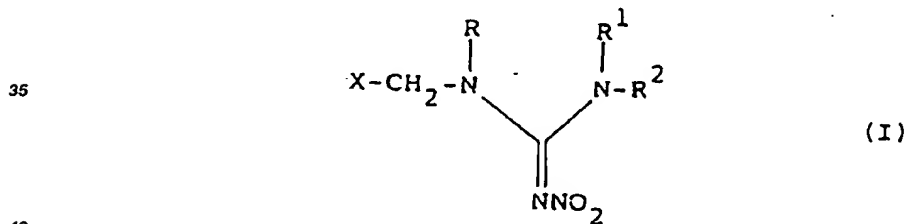
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As will be apparent from the foregoing, the compounds of this invention are extremely active for controlling various harmful insects and yet are low in toxicity to warm-blooded animals, fishes and Crustacea, the remaining quantity thereof after their use is small, and they do not produce phytotoxicity in various plants.

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### Claims

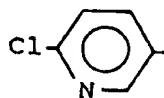
30 1. A nitroguanidine compound represented by the following general formula (I):



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wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by

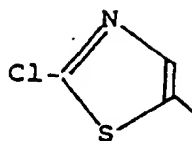
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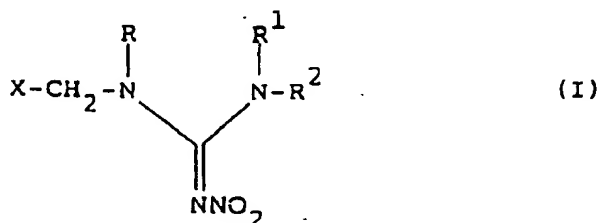
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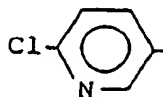
2. The nitroguanidine compound according to claim 1 which is N-[(6-chloro-3-pyridinyl)methyl]-N-methyl-N'-nitroguanidine.
- 10 3. The nitroguanidine compound according to claim 1 which is N-[(6-chloro-3-pyridinyl)methyl]-N,N',N'-trimethyl-N''-nitroguanidine.
4. The nitroguanidine compound according to claim 1 which is N-[(6-chloro-3-pyridinyl)methyl]-N'-methyl-N''-nitroguanidine.
- 15 5. The nitroguanidine compound according to claim 1 which is N-[(6-chloro-3-pyridinyl)methyl]-N'-nitroguanidine.
6. The nitroguanidine compound according to claim 1 which is N-[(6-chloro-3-pyridinyl)methyl]-N,N'-dimethyl-N''-nitroguanidine.
7. The nitroguanidine compound according to claim 1 which is N-[(2-chloro-5-thiazolyl)methyl]-N-methyl-N'-nitroguanidine.
- 20 8. The nitroguanidine compound according to claim 1 which is N-[(2-chloro-5-thiazolyl)methyl]-N,N',N'-trimethyl-N''-nitroguanidine.
9. The nitroguanidine compound according to claim 1 which is N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitroguanidine.
- 25 10. The nitroguanidine compound according to claim 1 which is N-[(2-chloro-5-thiazolyl)methyl]-N''-nitroguanidine.
11. A process for preparing a nitroguanidine compound represented by the following general formula (I):



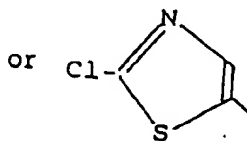
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40 wherein R stands for hydrogen or methyl; R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl; and X stands for a group indicated by



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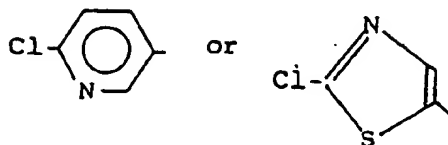
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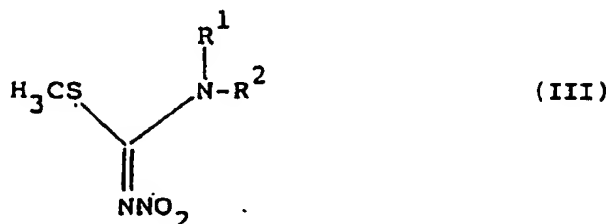
comprising the step of reacting, with or without solvents a compound represented by the general formula (II);



wherein R stands for hydrogen or methyl, and X stands for a group indicated by

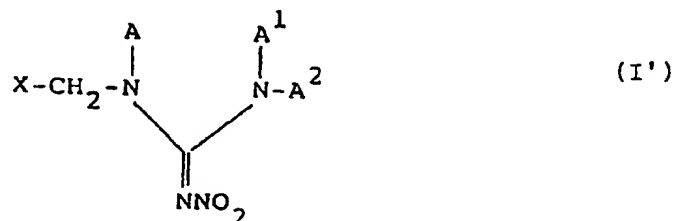


with a compound represented by the general formula (III):

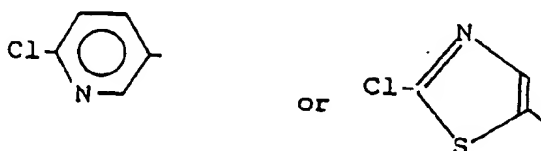


25 wherein R<sup>1</sup> and R<sup>2</sup> are the same or different, and stand for hydrogen or methyl.

12. A process for preparing a nitroguanidine compound represented by the following general formula (I'):

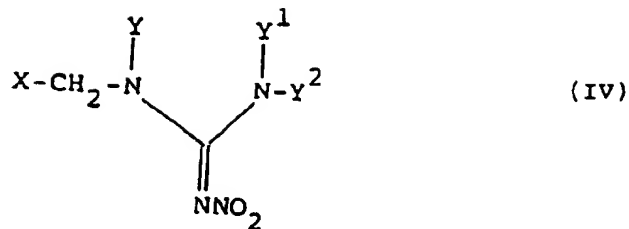


40 wherein A, A<sup>1</sup> and A<sup>2</sup> are the same or different, and stand for hydrogen or methyl provided that at least one of them is methyl, and X stands for a group indicated by

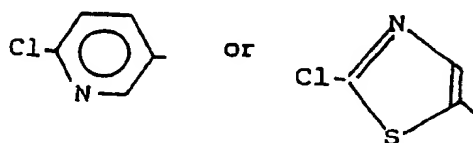


comprising the step of reacting, in the presence of a base a compound represented by the general formula (IV):

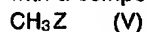
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wherein Y, Y<sup>1</sup> and Y<sup>2</sup> are the same or different, and stand for hydrogen or methyl provided that at least one of them is hydrogen; and X stands for a group indicated by



25 with a compound (V) represented by the general formula (V):



wherein Z stands for halogen.

13. An insecticide containing, a suitable carrier, and, as an active ingredient, a nitroguanidine compound represented by the general formula (1) according to any one of claims 1 to 10

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